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(71) Applicant (*for all designated States except US*): BIO-CHEMIE S.A. [ES/ES]; Ap. de Correos 97, E-08400 Granollers (ES).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): CENTELLAS, Victor [ES/ES]; Plaza Joan Alsina, 8,4-2-a, E-08440 Cardedeu (ES). GARCIA, Rafael [ES/ES]; Pza. Mercè Rodoreda, 3,3o,2a, E-08470 Sant Celoni (ES). POCH, Marta [ES/ES]; Av. Diagonal, 586,1o,1a, E-08028 Barcelona (ES). DIAGO, José [ES/ES]; Calle Josep Carner, 43,5-a, E-08400 Granollers (ES). LUDESCHER, Johannes [AT/AT]; Kleinsöll 101, A-6252 Breitenbach (AT). BOSCH, Immaculada [ES/ES]; Raval Cortines, 21, E-08500 Vic (ES).

(74) Agent: BECKER, Konrad; Novartis AG, Corporate Intellectual Property, Patent & Trademark Department, CH-4002 Basel (CH).

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(54) Title: MACROLIDES

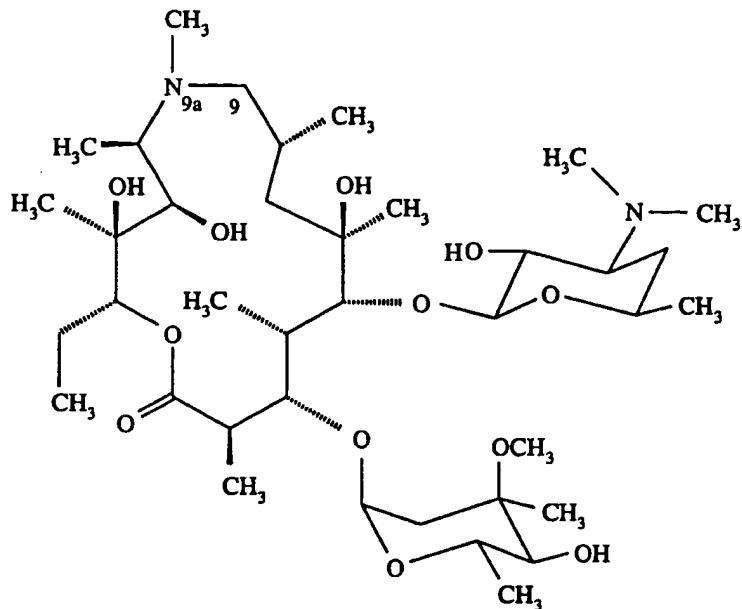
(57) Abstract: Azithromycin in the form of a stable monohydrate and processes for the preparation of azithromycin in the form of an, e.g. stable, monohydrate.

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Macrolides

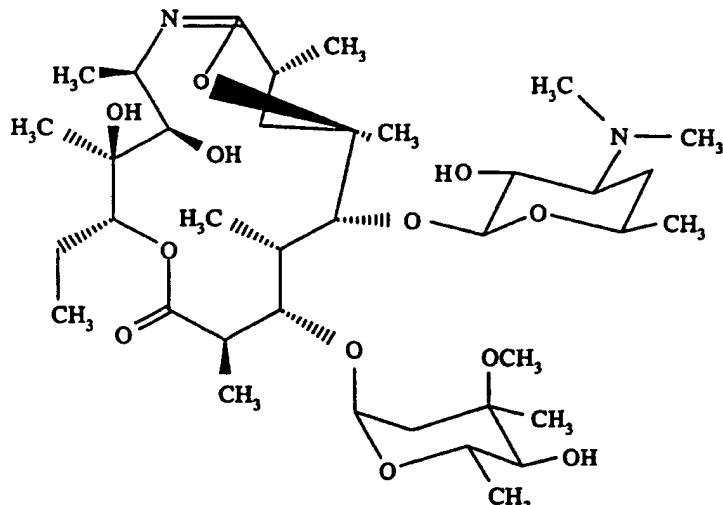
The present invention relates to macrolides, e.g. azithromycin and similar compounds.

Azithromycin (9-deoxo-9a-aza-9a-methyl-9a-homoerythromycin A) of formula I



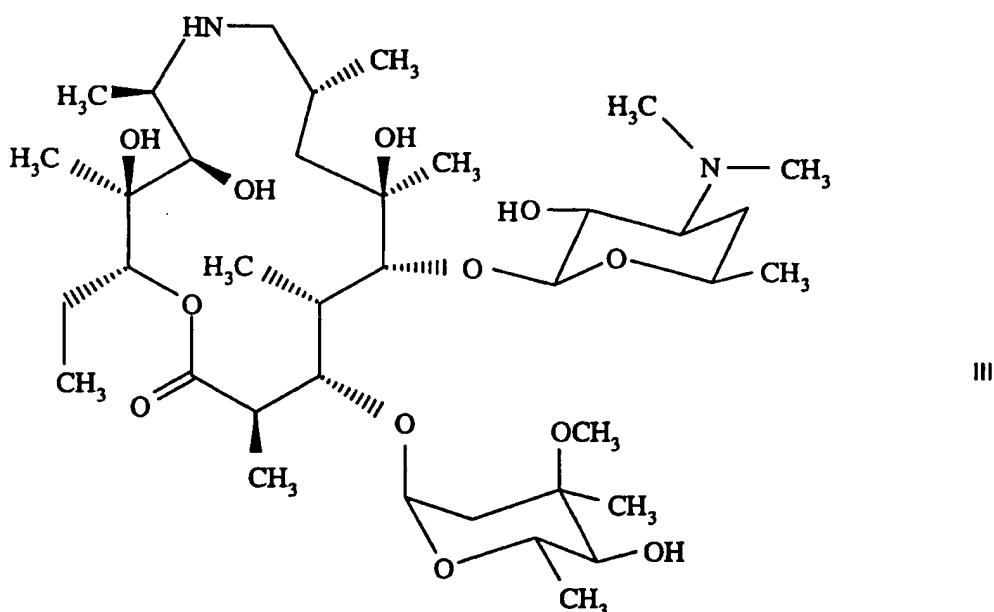
5 is a well-known antibacterial agent, described e.g. in The Merck Index, 12th edition (1996), page 157 (946) and may e.g. be produced by

- Beckmann rearrangement of erythromycin A 9 oxime to give 9-deoxo-6-deoxy-6,9-epoxy-9,9a-didehydro-9a-aza-9a-homoerythromycin A of formula

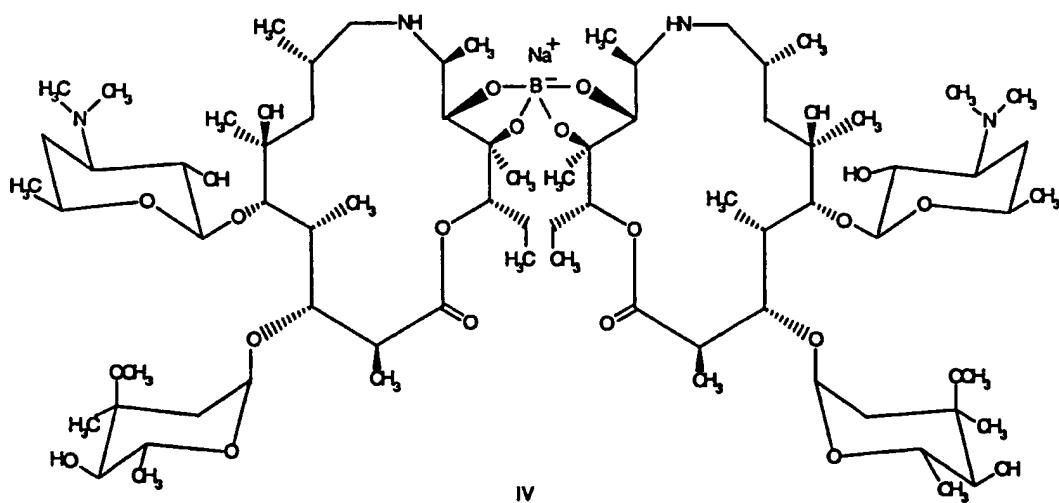


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- reduction of a compound of formula II to give 9-deoxo-9a-aza-9a-homoerythromycin A of formula



- N-methylation of a compound of formula III to give azithromycin, e.g. of formula I.
- Reduction of a compound of formula II to obtain a compound of formula III may be performed e.g. either catalytically under use of hydrogen; or, alternatively, in the presence of
- 5 borohydrides, e.g. sodium borohydride, which use, however, may result in intermediate borate compounds, e.g. esters of boric acid with one or more of the hydroxy groups of a compound of formula III. The chemical nature of resulting intermediate borate compounds may be dependent on reduction conditions used and different intermediate borate compounds and mixtures thereof may be obtained. E.g. according to data (MS, ^{11}B and ^{13}C NMR) e.g. an
- 10 intermediate borate compound of formula



may be obtained. Hydrolysis of intermediate borate compounds to obtain a compound of formula III may be carried out under acidic conditions which, however, may involve degrading conditions for the macrolide structure. Isolation of a compound of formula III from a reaction mixture resulting from intermediate borate compound-hydrolysis may be performed under

5 neutral or basic conditions and may be complicated because under neutral and basic conditions re-forming of intermediate boron compounds may occur and separation of a compound of formula III from intermediate borate compounds may e.g. afford chromatography. Thus, on industrial scale, reduction of a compound of formula II is preferably carried out catalytically under use of hydrogen.

10 A process was now found which surprisingly facilitates the isolation of a compound of formula III in a reaction mixture resulting from intermediate borate compound-hydrolysis and which may be carried out on technical scale.

15 In one aspect the present invention provides a process for the production of a compound of formula III, comprising the steps

- i) treating a compound of formula II with a boronated hydride;
- ii) hydrolyzing an intermediate borate compound formed in the presence of a polyhydroxylated compound; and, if desired

20 iii) isolating a compound of formula III from the reaction mixture.

In another aspect the present invention provides a process for the production of a compound of formula III, comprising the steps

- i) hydrolyzing an intermediate borate compound formed from a compound of formula II and

25 a boronated hydride in the presence of a polyhydroxylated compound; and, if desired,

- ii) isolating a compound of formula III from the reaction mixture.

It was surprisingly found that polyhydroxylated compounds present during hydrolysis of an intermediate borate compound may result in the formation of a borate ester, resulting from

30 reaction of boric acid which is generated during hydrolysis, and hydroxy groups of a polyhydroxylated compound. Thus, re-forming of intermediate borate compounds with the macrolide, e.g. a compound of formula III, may be suppressed or avoided and isolation of a compound of formula III may be facilitated.

35 A process according to the present invention may be carried out as follows:

A compound of formula II which is a known compound may be treated, e.g. stirred, in a solvent, e.g. dissolved, in the presence of boronated hydride, e.g. by addition of boronated hydride to a solution of a compound of formula II (or vice versa). A solution includes a suspension wherein a compound of formula II is at least in part dissolved. A solvent includes

5 solvent appropriate in reduction processes, preferably alcohols such as methanol, ethanol, isopropanol; and ethers, such as tetrahydrofuran; and mixtures of individual solvent, e.g. as cited above. An acid may be present, e.g. may be added to a solution of a compound of formula II, e.g. an acid which is appropriate to accelerate the reduction of imines, e.g. by formation of an iminium ion; including an organic acid, preferably formic acid or acetic acid,

10 and an inorganic acid, preferably hydrochloric acid or sulphuric acid; and mixtures of individual acids, e.g. as cited above. E.g. an acid in aqueous solution may be added to a solution of a compound of formula II. "Boronated hydride" includes a compound containing boron and hydrogen atoms, appropriate to act as a reducing agent, such as

15 - boranes or diboranes, e.g. in the form of stable complexes, such as a complex with tetrahydrofuran, methyl sulfide, pyridine, morpholine, 4-methylmorpholine, 1,4-oxathiane; borane complexes with amines, e.g. including ammonia, tert-butylamine, N,N-diethylaniline, N,N-diisopropylethylamine, dimethylamine, (4-dimethylamino)-pyridine, 4-ethylmorpholine, 2,6-lutidine, triethylamine, trimethylamine; and borane complexes with phosphines e.g. with diphenylphosphine, tributylphosphine, triphenylphosphine;

20 - metal borohydrides, such as sodium and potassium borohydride, sodium and potassium cyanoborohydride and lithium borohydride;

- alkylborohydrides, e.g. trialkylborohydrides, e.g. tri(C₁₋₄)alkyl borohydrides, such as lithium or sodium trimethyl- and triethylborohydride;

- alkoxyborohydrides, e.g. trialkoxyborohydrides, such as tri(C₁₋₄)alkoxyboro-hydrides, such as sodium trimethoxy-borohydride;

25 - acyloxyborohydrides, e.g. triacycloxyborohydrides, wherein acyl includes (C₂₋₆)acyl, such as sodium triacetoxyborohydride;

preferably sodium, potassium and lithium borohydride, sodium and lithium cyanoborohydride; and mixtures of individual boronated hydrides, e.g. as cited above;

30 e.g. in any appropriate form, such as in solid, e.g. powder, pellet, granule form; in solution, e.g. in 2-methoxyethylether, triethylene glycol dimethyl ether, aqueous solution, e.g. sodium hydroxide solution; e.g. fixed onto a carrier, such as alumina.

An intermediate borate compound, e.g. a borate ester, e.g. such as described above, e.g. of formula IV, may be formed in the reaction mixture and may be isolated, e.g. as conventional, e.g. by evaporating the solvent (system); or e.g. by hydrolizing an excess of boronated hydride

in the reaction mixture, e.g. by treating the reaction mixture with aqueous, e.g. inorganic, acid, adjusting the pH of the mixture obtained to a basic pH; and isolating an intermediate borate compound which may precipitate, e.g. by filtration; or extracting an intermediate borate compound into a solvent which is able to form a two-phase system with water and
5 which is able to dissolve an intermediate borate compound in a two-phase system with water; such as a halogenated hydrocarbon, e.g. methylene chloride; and, if desired evaporating off the solvent.

The amount of a boronated hydrate in respect with a compound of formula II is not critical. An appropriate amount may be easily determined by pre-testing.
10 An, e.g. isolated, intermediate borate compound obtained may be hydrolyzed, e.g. in a solvent, e.g. in solution, such as an aqueous solution, e.g. water or an organic solvent which is inert under the reaction conditions and which contains water, e.g. in the presence of an acid, e.g. an inorganic acid, such as hydrochloric acid, sulphuric acid and in the presence of a polyhydroxylated compound, e.g. by addition of a polyhydroxylated compound to a solution
15 (including a suspension) of an intermediate borate compound. An intermediate borate compound obtained in a solution of a solvent which forms a two-phase system with water may e.g. be extracted into an acidic aqueous solvent and a polyhydroxylated compound may be added to the acidic mixture obtained.

A polyhydroxylated compound includes an organic compound having two or more hydroxyl
20 groups, e.g.

- linear polyalcohols, such as ethyleneglycol, propyleneglycol, glycerol, 1,2-hexanediol, 1,6-hexanediol, 2,5 hexanediol, 1,2,3-hexanetriol, 1,2,6-hexanetriol or pentaerythritol; linear polyalcohols, e.g. as cited above, having beside at least two hydroxyl groups further functional groups, e.g. amine group(s), such as diethanolamine or triethanolamine;
- cyclic polyalcohols; e.g. exo-cyclic polyalcohols such as exo-2,3-norbornanediol;
- sugar-type polyalcohols, e.g. monosaccharides, such as glyceraldehyde, erythrose, erythrulose, arabinose, lyxose, ribose, ribulose, xylose, xylulose, allose, altrose, fructose, galactose, glucose, mannose, sorbose, tagatose and talose, e.g. in any of its isomeric forms (D-form, L-form or DL form);
- sugar-type polyalcohols obtainable by reduction of corresponding sugars, such as erythrol, threitol, arabitol, xylitol, adonitol, sorbitol, dulcitol, manitol;
- sugar-type polyalcohols having beside at least two hydroxy groups further functional groups, e.g. amines such as glucamine or N-methylglucamine;
- a resin, e.g. an ion exchange resin, containing hydroxy groups, e.g. Amberlite IRA-743 resin, e.g. as offered in the Catalogue Handbook of Fine Chemicals (1996-1997),

in any appropriate form, e.g. in solid form, in solution, fixed onto a carrier.

The amount of a polyhydroxy compound in respect with an intermediate borate compound is not critical. An appropriate amount may be easily determined by pre-testing.

A compound of formula III may be obtained and may be isolated from the reaction mixture,

5 e.g. as conventional, e.g. by filtration; or e.g. may be extracted into a solvent which is able to form a two-phase system with water and which is able to dissolve a compound of formula III in a two-phase system with water; e.g. a two-phase system may be obtained, the phases may be separated and the solvent of the organic phase may be removed, e.g. by evaporation, e.g. to dryness; or a solution of a compound of formula III may be used as such, e.g. without further
10 isolation or purification, in a methylation reaction, e.g. to obtain azithromycin.

In a further aspect the present invention provides a process for the production of

azithromycin, e.g. of formula I, e.g. in the form of a solvate, comprising

i) treating a compound of formula II with a boronated hydride,
15 ii) hydrolyzing an intermediate borate compound formed in the presence of a polyhydroxylated compound to obtain a compound of formula III, and
iii) methylating a compound of formula III obtained in step ii) at the amine group in position 9a of the erythromycin ring structure; and, if desired,
iv) isolating azithromycin; e.g. in the form of a solvate, e.g. in the form of a hydrate, such
20 as a dihydrate or a monohydrate.

In another aspect the present invention provides a process for the production of azithromycin,

e.g. of formula I, e.g. in the form of a solvate, comprising the steps

i) hydrolyzing an intermediate borate compound formed from a compound of formula II
25 and a boronated hydride in the presence of a polyhydroxylated compound;
ii) methylating a compound of formula III obtained in step ii) at the amine group in position 9a of the erythromycin ring structure; and
iii) isolating azithromycin; e.g. in the form of a solvate, e.g. in the form of a hydrate, such as
a dihydrate or a monohydrate.
30

In a further aspect the present invention provides the use of a compound of formula III

obtained by hydrolyzing an intermediate borate compound formed from a compound of

formula II and a boronated hydride, in the presence of a polyhydroxylated compound

in the production of azithromycin, e.g. in the form of a solvate, e.g. in the form of a hydrate,

35 such as a dihydrate or a monohydrate.

Methylation of a compound of formula III at the amine group in position 9a of the erythromycin ring structure may be carried out as appropriate, e.g. according to a method as conventional, e.g. treating a solution of a compound of formula III in a solvent, e.g. ethyl acetate, with formic acid and aqueous formaldehyde and isolating azithromycin obtained.

Processes according to the present invention are useful in the production of azithromycin. Advantages include e.g. high yields in the production of desired compounds, e.g. of formula III and of azithromycin; a compound of formula III may be obtained in pure form and may be used without further purification or even without isolation in a subsequent reaction, e.g. in a methylation step; the processes may be used on industrial scale.

Azithromycin, e.g. obtainable according to a process of the present invention, may be in the form of a solvate, e.g. in the form of a hydrate, such as a monohydrate or e.g. in the form of a dihydrate. It is known that azithromycin in the form of a monohydrate is unstable, e.g. the crystall structure of azithromycin in the form of a monohydrate may break down under normal air humidity conditions within some hours. Thus, azithromycin in the form of a monohydrate which may be obtained according to a conventional method, e.g. by precipitation with water from ethanolic solution, is described to be not easy to handle. This might be one reason why azithromycin currently on the market is in the form of a dihydrate which is known to be stable under normal air humidity conditions, e.g. the crystall structure of azithromycin in the form of a dihydrate is known not to break down within some hours under normal air humidity conditions.

Azithromycin in the form of a monohydrate produced according to known methods, e.g. precipitation with water from an ethanolic solution, may beside its instability contain a high content of residual solvents, e.g. 1.0% and more and may be inappropriate for use as a pharmaceutical.

It was now surprisingly found that azithromycin, e.g. of formula I, may be obtained in the form of an, e.g. crystalline, monohydrate which is stable and which may be easy to handle. "Stable monohydrate" as used herein includes a compound of formula I in the form of a monohydrate which is crystalline and which maintains its crystalline structure for at least 24 hours, e.g. for several weeks, under normal, e.g. normal air humidity, conditions. The crystalline structure of azithromycin in the form of a monohydrate may be determined by its (known) X-ray powder diffraction pattern.

In another aspect the present invention provides azithromycin, e.g. of formula I, in the form of a stable, e.g. crystalline, monohydrate.

5 It was also found that azithromycin in the form of a stable monohydrate according to the present invention may contain water from, e.g. about, 1.5%, e.g. and more, such as 2.0%; up to 5.5% and more; e.g. 6.0% w/w, e.g. 2.0% to 6.0% w/w; such as 1.5% to 5.5% w/w without changing its crystalline structure and without taking up remarkably further water. A water content of 2.0% to 6.0% w/w includes the water necessary for azithromycin
10 monohydrate formation.

In another aspect the present invention provides a composition comprising, e.g. consisting essentially of; azithromycin, e.g. of formula I, in the form of a stable monohydrate, e.g. crystalline; e.g. showing an X-ray powder diffraction pattern corresponding to azithromycin in
15 the form of a monohydrate, e.g. even after 24 hours under normal air humidity conditions; and 2.0% to 6.0% w/w of water.

It was also found that azithromycin in the form of a stable, e.g. crystalline, monohydrate may be produced by removing organic solvent from a solution of azithromycin in organic solvent;
20 e.g. whereas azithromycin produced according to prior art, e.g. by precipitation with water from ethanolic solution may result in azithromycin in the form of an unstable monohydrate.

In another aspect the present invention provides a process for the production of azithromycin in the form of a stable monohydrate, e.g. crystalline, comprising removing organic solvent
25 from a solution of azithromycin in organic solvent.

A process according to the present invention may be carried out as follows:

Azithromycin in any form, e.g. in free base form; and in the form of a salt, e.g. in the form of a hydrochloride, e.g. a dihydrochloride, acetate; and/or in the form of a solvate, e.g. in the
30 form of an (unstable) monohydrate or in the form of a dihydrate; and in non-solvate form, e.g. anhydrous form, may be used as a starting material and may be dissolved, e.g. suspended, in organic solvent. A solution of azithromycin in organic solvent may be obtained. A "solution" includes a suspension, in which at least a part of azithromycin is dissolved. Organic solvent includes solvent which is appropriate to dissolve azithromycin, preferably water-miscible
35 organic solvent. Water-miscible organic solvent includes organic solvent which may form a

one-phase system with water, preferably water-miscible ketones such as acetone; lower alcohols, e.g. (C_{1-4})alcohols, such as methanol, ethanol, isopropanol; lower alkyl acetates, such as methyl acetate, ethyl acetate; and acetic acid and formic acid amides, such as dimethyl acetamide, N,N-dimethylformamide; or a mixture of individual organic solvents, e.g. as listed above. Water may be present in organic solvent. An appropriate temperature for the production of a solution of azithromycin in organic solvent includes a temperature at which azithromycin is not degraded, including, e.g. a temperature range of -20°C to 70 °C, such as 0°C to 50 °C. If as a starting material azithromycin is used in the form of a salt, a base may be used to provide azithromycin in solution in the form of a free base, e.g. to obtain an appropriate pH of a solution of azithromycin, e.g. an pH where azithromycin is in the form of a free base, e.g. by addition of a base to a mixture of azithromycin in the form of a salt with organic solvent, e.g. according to a method as conventional. The concentration of a solution of azithromycin in organic solvent is not critical; preferably a saturated or almost saturated solution may be used. An amount of water which is necessary for the formation of azithromycin in the form of a monohydrate may be present in the organic solvent. If a solution of azithromycin in organic solvent contains less water than necessary for azithromycin monohydrate formation, water may be added, e.g. sufficient water to form azithromycin monohydrate. If a solution of azithromycin in organic solvent contains more water, e.g. an excess of water, than necessary for azithromycin monohydrate formation, excess of water may be removed, if desired, but not necessarily. Appropriate methods for removal of excess of water includes methods as conventional, e.g. removing, e.g. distilling off, organic solvent, e.g. preferably lower alcohols, lower alkyl acetates or acetic and formic acid amides, together with water from a solution of azithromycin in organic solvent; or addition of a drying agent, e.g. such as conventional in organic chemistry, e.g. sodium sulphate, prior to organic solvent removal of a solution of azithromycin in organic solvent, preferably in lower ketones. Organic solvent from a solution of azithromycin in organic solvent may be removed, e.g. partially or (almost) completely, e.g. by a method as conventional, e.g. by distillation, such as evaporation.

Alternatively, azithromycin in the form of a monohydrate having a water content which is at least in the range of azithromycin in the form of a monohydrate, or higher, e.g. unstable azithromycin in the form of a monohydrate, or azithromycin in the form of a dihydrate may be warmed up to an appropriate temperature, e.g. as described above, in organic solvent, e.g. in lower alcohols, to form a solution of azithromycin in alcohol; and the solution may be cooled to obtain azithromycin, e.g. crystalline, in the form of a stable monohydrate. In order

to increase yields alcohol may be removed from a correspondingly obtained crystal suspension of azithromycin in the form of a stable monohydrate in alcohol.

Azithromycin in the form of a stable monohydrate may precipitate, e.g. in crystalline form and may be isolated, e.g. as conventional, e.g. by filtration or centrifugation. An anti-solvent with
5 the exception of water, e.g. an organic solvent which decreases the solubility of azithromycin in organic solvent solution, may be added to a residue obtained by organic solvent removal in order to increase yields.

In another aspect the present invention provides a process for the production of azithromycin
10 in the form of a stable monohydrate comprising dissolving azithromycin in an alcohol, and crystallizing azithromycin in the form of a stable monohydrate from the alcoholic solution.

In another aspect the present invention provides a process for the production of azithromycin
15 in the form of a stable monohydrate comprising dissolving azithromycin in the form of a dihydrate in organic solvent, e.g. ethanol, isopropanol, methyl acetate; e.g. or a or a mixture of individual solvents listed, distilling off water-miscible organic solvent and, if desired, e.g. and if present, excess of water; and isolating azithromycin in the form of a stable monohydrate.

20 In another aspect the present invention provides a process for the production of azithromycin in the form of a stable monohydrate comprising dissolving azithromycin in the form of a dihydrate in organic solvent, e.g. acetone, e.g., if desired drying the solution obtained, e.g. by addition of a drying agent; distilling off water-miscible organic solvent and isolating azithromycin in the form of a stable monohydrate.

25 In a further aspect, the present invention provides a composition containing, e.g. consisting essentially of; azithromycin in the form of a monohydrate, e.g. solid, e.g. crystalline, e.g. stable; and methyl acetate, acetone or isopropanol.

30 It was also surprisingly found that azithromycin, e.g. of formula I, may be obtained in the form of an, e.g. crystalline, monohydrate, e.g. which is stable, and which may be easy to handle; and which may have a low content of (residual) organic solvent, e.g. a content of organic solvent, e.g. including organic solvent whith low toxic potential (e.g. Class 3 solvents); of 0.5% w/w and lower; e.g. amounts which are on the lower end of analytical detectability,
35 such as 0.001%, e.g. 0.002%, e.g. 0.003%, e.g. 0.1% to 0.5%; e.g. and less; e.g. fulfilling

European Pharmacopoeia specifications. E.g. azithromycin in the form of a monohydrate, e.g. stable; e.g. crystalline, having a low organic solvent content in an amount of the lower end of analytical detectability up to 0.5%; e.g. and even lower; may be e.g. obtained as described above, or, e.g. more specifically, as follows:

5 Azithromycin in any form, e.g. in a form as described above as a starting material for the production of azithromycin in the form of a stable monohydrate, may be used as a starting material. A solution of azithromycin in the form of a salt in a solvent may be produced, e.g. either by dissolving azithromycin in form of a salt in a solvent; or by conversion of azithromycin in free form in a solvent into the form of a salt of azithromycin; e.g. by addition 10 of an acid to azithromycin in solvent. A "solution" includes a suspension, in which at least a part of azithromycin in the form of a salt is dissolved. Appropriate acids include, e.g. an organic acid, for example formic acid or acetic acid, and an inorganic acid, for example hydrochloric, hydrobromic, nitric or sulphuric acid, preferably hydrochloric acid or sulphuric acid. Solvent includes solvent which is appropriate to dissolve azithromycin in the form of a 15 salt, e.g. including aqueous solvent. Aqueous solvent includes water or a mixture of water with organic solvent, e.g. one or more organic solvents, for example water miscible and water immiscible organic solvent, such as alcohols, e.g. methanol, ethanol, isopropanol; ketones such as acetone, methyl isobutyl ketone; alkyl esters such as alkyl esters, e.g. (C₁₋₄)alkyl; of formic or acetic acid, e.g. methyl acetate, ethyl acetate, isopropyl acetate, butyl acetate; aromatic 20 hydrocarbons such as toluene, xylenes; ethers, such as tetrahydrofuran, methyl t-butyl ether; chlorinated hydrocarbons such as methylene chloride; and amides such as monoalkyl and dialkyl amides, e.g. N-methyl formamide, N,N-dimethylacetamide, N,N-dimethylformamide; preferably water or a mixture of water with one or more alcohols, ketones, alkyl acetates; e.g. water, or aqueous solvent, such as water or water containing 0.5% to 20 % v/v; such as 1% 25 to 15 % v/v of organic solvent.

Appropriate reaction conditions for the production of a solution of azithromycin in the form of a salt according to the present invention include, e.g.

(i) A temperature at which azithromycin is not degraded, including, e.g. a temperature range of -20 °C to 90 °C , such as 0°C to 70 °C;

30 (ii) An appropriate pressure, e.g. atmospheric pressure, and a pressure which is above or below atmospheric pressure;

(iii) Appropriate dilution, e.g. a dilution range of 1 g to 500 g of azithromycin in the form used as a starting material, per litre of solvent.

A resulting solution of azithromycin in the form of a salt in a solvent may be purified as 35 appropriate, e.g. by filtration, charcoal treatment; in order to remove impurities. The pH of

an, e.g. purified, solution of azithromycin in the form of a salt may be adjusted to an pH where azithromycin is present in free form, including e.g. a pH of, e.g. ca., 8.0 to 13.0, such as 9.0 to 12.0, e.g. 10.0 to 11.0; e.g. by addition of a base to a solution of azithromycin in the form of a salt in a solvent. A "solution" of azithromycin in free form includes a suspension, in which at least a part of azithromycin in free form is dissolved. Appropriate bases include bases which are suitable for pH adjustment, e.g. an inorganic base, such as ammonia or an alkali-, e.g. sodium, potassium; earth alkali-, e.g. calcium, magnesium; and ammonium-; -hydroxide, -carbonate, -hydrogencarbonate; and an organic base, such as an amine, e.g. an alkyl amine, or a mixture of individual bases, e.g. as described above. A base may be preferably a hydroxide, e.g. sodium or ammonia; preferably in aqueous solution.

Azithromycin in free form and in the form of an, e.g. stable; monohydrate, e.g. crystalline; may precipitate from the solution and may be isolated, e.g. according to a method as conventional, e.g. by centrifugation or filtration, and may be dried, for example at appropriate temperatures, e.g. including a range of, e.g. ca., 30°C to 80 °C. Azithromycin, e.g. crystalline, in the form of an, e.g. stable; monohydrate may be obtained; e.g. a composition of azithromycin in the form of an, e.g. stable, monohydrate and less than 0.5% w/w of organic solvent; e.g. a composition comprising, e.g. consisting essentially of; azithromycin in the form of a stable monohydrate and more than 0% to 0.5% w/w of organic solvent; e.g. analytically detectable amounts of organic solvent up to 0.5% w/w; e.g. and containing 2.0% to 6.0% w/w of water.

In another aspect the present invention provides a process for the production of azithromycin in the form of an, e.g. stable, monohydrate, e.g. crystalline, comprising the steps of i) adjusting the pH of a solution of azithromycin in the form of a salt, and ii) isolating azithromycin of formula I in the form of an, e.g. stable, monohydrate, characterised in that the solution of azithromycin in step i) is an aqueous solution wherein the solvent is selected from water or a mixture of water and organic solvent.

In another aspect the present invention provides a composition comprising, e.g. consisting essentially of; azithromycin, e.g. of formula I, in the form of an, e.g. crystalline, monohydrate, e.g. stable; and organic solvent in an amount of 0.5% w/w and lower; e.g. and an amount which is higher than 0% w/w; e.g. an analytically detectable amount up to 0.5% w/w; e.g. an amount which fulfils European Pharmacopoeia specifications.

When solely water is used as a solvent according to the present invention, azithromycin in the form of an, e.g. stable, monohydrate, e.g. crystalline, may be obtained, substantially free of organic solvent.

5 In another aspect the present invention provides azithromycin in the form of a monohydrate, substantially free of organic solvent.

Azithromycin in the form of a stable monohydrate according to the present invention; e.g. and a composition of azithromycin in the form of a monohydrate, e.g. stable, and an analytically 10 detectable amount up to 0.5% w/w of organic solvent; and azithromycin in the form of an, e.g. stable, monohydrate substantially free of organic solvent; is useful in the production of a pharmaceutical composition containing azithromycin as an active ingredient.

In another aspect the present invention provides a pharmaceutical composition, comprising, 15 e.g. essentially consisting of,

- azithromycin in the form of a stable monohydrate; or
- a composition of azithromycin in the form of a monohydrate, e.g. stable, and an analytically detectable amount up to 0.5% w/w of organic solvent; or
- azithromycin in the form of an, e.g. stable, monohydrate substantially free of organic 20 solvent;

in association with at least one pharmaceutical carrier or diluent.

A pharmaceutical composition comprising, e.g. essentially consisting of,

- azithromycin in the form of a stable monohydrate as an active ingredient; or
- a composition of azithromycin in the form of a monohydrate, e.g. stable, and an analytically detectable amount up to 0.5% w/w of organic solvent; or
- azithromycin in the form of an, e.g. stable, monohydrate substantially free of organic solvent;

according to the present invention may contain the same concentrations of azithromycin and 30 may be used for the same indications in the same dosage ranges as a known pharmaceutical composition containing azithromycin in the form of a dihydrate as an active ingredient, e.g. as is currently on the market.

The following examples illustrate the present invention. All temperatures are given in degree 35 Celsius and are uncorrected.

The X-ray powder diffraction pattern and the IR spectrum of azithromycin in the form of a monohydrate obtained according to the following examples corresponds to that of known (unstable) azithromycin in the form of a monohydrate.

Azithromycin in the form of a monohydrate obtained according to the following examples 5 maintains its crystallinity and its X-ray powder diffraction pattern when kept for several weeks under normal air humidity conditions.

Water content (% w/w) is determined by the K.Fischer method and includes the water necessary for azithromycin monohydrate formation where indicated.

Residual solvent (% w/w) is determined by GC, head-space method.

Example 1

To a solution of 35.62 g of a compound of formula II (assay 92.8%) in 180 ml of methanol, cooled to -10/-15 °, 14.25 g of sodium borohydride are added during ca. 2 hours. The reaction mixture obtained is kept at -10/-15° for ca. further 2 hours, heated to 55°, cooled to 20° and the solvent is evaporated off. The residue obtained (108.9 g) is dissolved in a mixture of methylene chloride and water and is stirred at room temperature for ca. 30 minutes. A two-phase system is formed. The organic phase is separated off and the aqueous phase is extracted with methylene chloride. The organic phase is dried over sodium sulphate and the solvent is evaporated off.

31.15 g of a residue are obtained. The residue obtained is an intermediate borate compound and believed to be a compound of formula IV, e.g. confirmed by the following data:

Elemental analysis: Theory: B: 0.7%; Na: 1.0%. Found: B: 0.6%; Na: 1.1%

MS-FAB (-): 1476 ($M^+ - Na^+ + H^+$). An ^{11}B -NMR spectrum shows a ppm signal in the expected range of a tetracoordinated boron bond to four oxygen atoms..

Example 2

1.98 g of an intermediate borate compound obtained according to example 1 are suspended in 34 ml of water. 20% sulphuric acid is added to the suspension obtained and a pH of 2.8 is adjusted. 22 g of an ion resin containing hydroxy groups (Amberlite IRA-743) are added to the mixture obtained and the mixture obtained is stirred for ca. 30 minutes, the resin is filtrated off and washed with water. The pH of the filtrate obtained is adjusted to a basic pH with 20% sodium hydroxide. The mixture obtained is extracted with ethyl acetate, the organic phase is dried and evaporated off.

Yield: 1.22 g of a compound of formula III in pure form.

Example 3

Is carried out according to the method as described in example 2 but using 1.82 g of N-methyl-D-glucamine instead of 10.22 g of Amberlite IRA-743 resin.

Yield: 1.33 g of a compound of formula III.

Example 4

Is carried out according to the method as described in example 2 but using 10.02 g of an intermediate borate compound obtained according to example 1 instead of 1.98 g, 170 ml of

water instead of 34 ml and 8.60 g of sorbitol instead of 10.22 g of Amberlite IRA-743 resin.
Yield: 8.50 g of a compound of formula III.

Example 5

5 6.48 g of a compound of formula III obtained according to example 4 are dissolved in 58 ml of ethyl acetate and 0.6 ml of formic acid and 1.30 ml of 37% aqueous formaldehyde are added to the solution obtained. The mixture obtained is refluxed for ca. 2 hours. HPLC analysis shows the formation of azithromycin. Yield: 77%.of theory.

10 **Example 6**

70.0 g of azithromycin dihydrate are dissolved in 280 ml of methyl acetate at 25°. The resulting solution is heated to 55-60° and 225 ml of methyl acetate are distilled off. The distillation residue is cooled to 20°. Azithromycin in the form of a monohydrate crystallizes, is filtrated off and dried. Yield: 47.5 g. Water content: 2.5%

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Example 7

15.0 g of azithromycin dihydrate are dissolved in 75 ml of acetone, anhydrous sodium sulphate is added, the mixture obtained is stirred for ca. 10 minutes and the solid is filtrated off and washed with 15 ml of acetone. From the solution obtained 57 ml of acetone are
20 distilled off and the resulting suspension is cooled to room temperature. Azithromycin in the form of a monohydrate crystallizes, is filtrated off and dried.

Yield: 7.3 g. Water content: 2.6%

Example 8

25 10.0 g of azithromycin dihydrate are suspended in 10 ml of absolute ethanol and the mixture is heated. A clear solution is obtained, which is cooled slowly to room temperature and stirred at ca. 25° for ca. one hour. Azithromycin in the form of a monohydrate crystallizes. Ethanol is evaporated off and azithromycin in the form of a monohydrate is filtrated off and dried.
Yield: 5.26 g. Water content: 1.9%

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Example 9

Example 8 is repeated but using 10 ml of isopropanol instead of 10 ml of absolute ethanol.
Crystalline azithromycin in the form of a monohydrate is obtained.
Yield: 6.0 g, Water content: 2.5%.

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Example 10

20 g of azithromycin in the form of a dihydrate, suspended in 40 ml of water are treated with 2N HCl. A solution is obtained, is filtered and the filtrate is added dropwise to 80 ml of water keeping the pH between 10 and 11 by addition of 0.5 N sodium hydroxide at a temperature of 5 ca. 55°. The suspension obtained is cooled to room temperature and stirred for ca. 30 minutes. A solid precipitate obtained is filtrated off, washed with water and dried. 18.2 g of azithromycin in the form of a monohydrate are obtained. Water content: 3.7%. Residual solvent: acetone 0.01%.

Azithromycin assay on anhydrous basis (HPLC): 98.6%.

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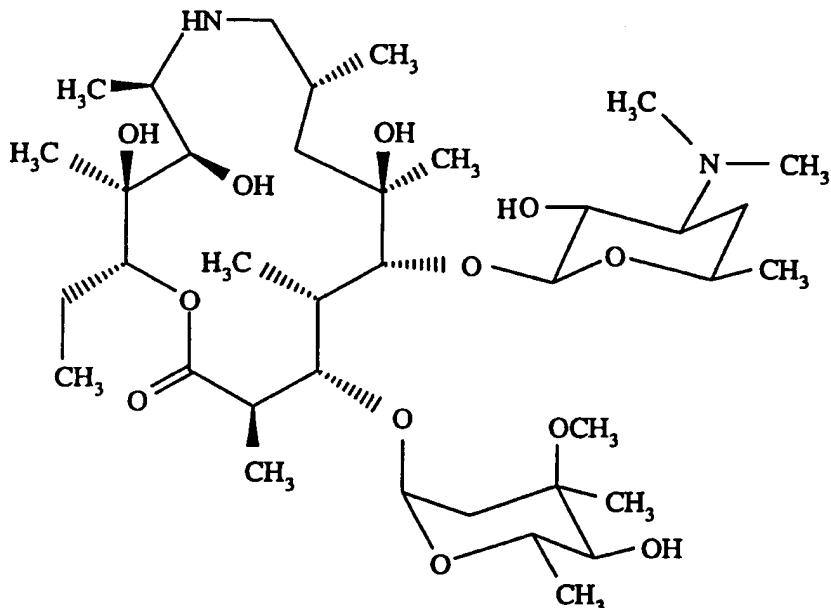
Example 11

Example 10 is repeated, but using a mixture of 80 ml of water and 4 ml of ethanol instead of 80 ml of water in the precipitation step; 19.5 g of azithromycin in the form of a monohydrate are obtained. Water content: 2.9%.

15 Residual solvents: acetone 0.003%, ethanol 0.002%.

Patent claims

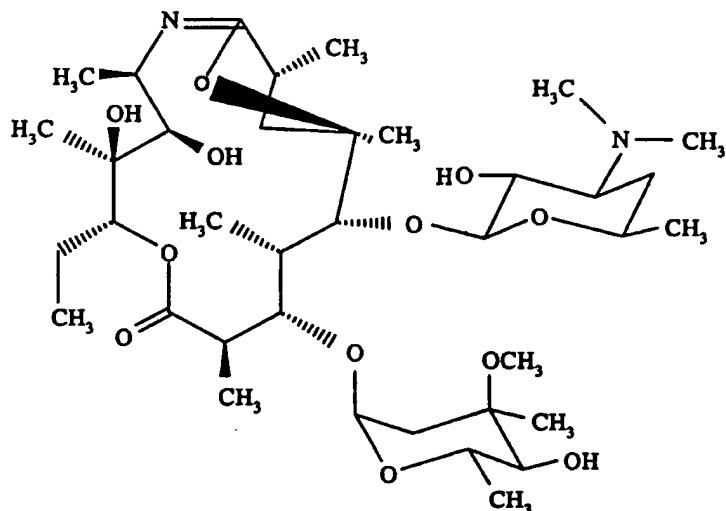
1. A process for the production of a compound of formula



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comprising the steps

i) treating a compound of formula



II

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with a boronated hydride,

ii) hydrolyzing an intermediate borate compound formed in the presence of a polyhydroxylated compound; and, if desired,

iii) isolating a compound of formula III from the reaction mixture.

2. A process for the production of a compound of formula III as defined in claim 1, comprising the steps

5 i) hydrolyzing an intermediate borate compound formed from a compound of formula II as defined in claim 1 and a boronated hydride in the presence of a polyhydroxylated compound; and, if desired

ii) isolating a compound of formula III from the reaction mixture.

10 3. A process for the production of azithromycin, comprising

 i) treating a compound of formula II as defined in claim 1, with a boronated hydride,

 ii) hydrolyzing an intermediate borate compound formed in the presence of a polyhydroxylated compound to obtain a compound of formula III as defined in claim 1,

15 iii) methylating a compound of formula III obtained in step ii) at the amine group in position 9a of the erythromycin ring structure; and, if desired,

 iv) isolating azithromycin; e.g. in the form of a solvate.

20 4. A process for the production of azithromycin comprising the steps

 i) hydrolyzing an intermediate borate compound formed from a compound of formula II, as defined in claim 1, and a boronated hydride in the presence of a polyhydroxylated compound;

 ii) methylating a compound of formula III as defined in claim 1, at the amine group in position 9a of the erythromycin ring structure; and

25 iii) isolating azithromycin.

25 5. A process according to any one of claims 4 or 5, comprising that azithromycin is isolated in the form of a solvate.

30 6. Use of a compound of formula III obtained by hydrolyzing an intermediate borate compound formed from a compound of formula II, as defined in claim 1, and a boronated hydride in the presence of a polyhydroxylated compound; in the production of azithromycin.

35 7. Azithromycin in the form of a stable monohydrate.

8. A composition comprising azithromycin in the form of a stable monohydrate; and 2.0% to 6.0% w/w of water.
- 5 9. A process for the production of azithromycin in the form of a stable monohydrate, comprising removing organic solvent from a solution of azithromycin in organic solvent.
10. A process for the production of azithromycin in the form of a stable monohydrate comprising dissolving azithromycin in an alcohol, and crystallizing azithromycin in the form of a stable monohydrate from the alcoholic solution.
- 15 11. A process for the production of azithromycin in the form of a stable monohydrate comprising dissolving azithromycin in the form of a dihydrate in ethanol, isopropanol or methyl acetate, distilling off organic solvent; and, if desired excess of water; and isolating azithromycin in the form of a stable monohydrate.
- 20 12. A process for the production of azithromycin in the form of a stable monohydrate comprising dissolving azithromycin in the form of a dihydrate in acetone, distilling off water-miscible organic solvent and isolating azithromycin in the form of a stable monohydrate.
13. A composition comprising azithromycin in the form of a monohydrate and methyl acetate, acetone or isopropanol.
- 25 14. A process for the production of azithromycin in the form of a monohydrate comprising the steps of
 - i) adjusting the pH of a solution of azithromycin in the form of a salt, and
 - ii) isolating azithromycin of formula I in the form of a monohydrate, characterised in that the solution of azithromycin in step i) is an aqueous solution wherein the solvent is selected from water or a mixture of water and organic solvent.
- 30 15. A composition comprising azithromycin in the form of a monohydrate and organic solvent in an analytically detectable amount up to 0.5% w/w.
- 35 16. Azithromycin in the form of a monohydrate, substantially free of organic solvent.

17. A pharmaceutical composition comprising

- azithromycin in the form of a stable monohydrate; or
- a composition of azithromycin in the form of a monohydrate and an analytically detectable amount up to 0.5% w/w of organic solvent; or
- azithromycin in the form of a monohydrate substantially free of organic solvent; in association with at least one pharmaceutical carrier or diluent.

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18. Azithromycin in the form of a monohydrate according to any one of claims 7, 8, 13, 15, 10 16 and 17; or obtained by a process according to any one of claims 9, 10, 11, 12 or 14; which is crystalline.

19. Azithromycin in the form of a monohydrate according to any one of claims 13, 15 or 16; or obtained by a process according to claim 14; which is a stable monohydrate.

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 00/05948

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07H17/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07H

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data, BIOSIS, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	BAYOD-JASANADA M ET AL: "SYNTHESIS OF 9-DEOXO-9A-AZA-HOMOERYTHROMYCIN A 11,12-HYDROGEN BORATE AND AZITHROMYCIN 11,12-HYDROGEN BORATE. A NEW PROCEDURE TO OBTAIN AZITHROMYCIN DIHYDRATE" JOURNAL OF ORGANIC CHEMISTRY, US, AMERICAN CHEMICAL SOCIETY, EASTON, vol. 62, no. 21, 1997, pages 7479-7481, XP002061020 ISSN: 0022-3263 page 7479, scheme 1 page 7450, line 1	1-6
X	----- -/-	7,8,13, 15-19

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

20 October 2000

Date of mailing of the international search report

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Scott, J

INTERNATIONAL SEARCH REPORT

International Application No PCT/EP 00/05948	
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 298 650 A (PFIZER) 11 January 1989 (1989-01-11) page 4-5, preparation 1 page 4, preparation 1	1-6
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A	WO 98 41532 A (BIOCHEMIE SA ;DIAGO JOSE (ES); BOSCH IMMACULADA (ES); CENTELLAS VI) 24 September 1998 (1998-09-24) page 6, line 19 -page 7, line 8; claims 1-10; examples 1-3	1-6
P,X	EP 0 984 020 A (APOTEX INC) 8 March 2000 (2000-03-08) claims 1-9	7-11, 13-19
P,X	WO 00 27856 A (BIOCHEMIE SA ;DIAGO JOSE (ES); BOSCH IMMACULADA (ES); CENTELLAS VI) 18 May 2000 (2000-05-18) page 6, lines 5-7; page 8; Claims 1-8	7-10, 13-19
P,X	EP 0 941 999 A (HOVIONE INT LTD) 15 September 1999 (1999-09-15) example 4	7-10, 13-19

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/EP 00/05948**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-6

Process for preparing a compound of formula III, claim 1 or process for preparing azithromycin using a boronated hydride and a polyhydroxylated compound.

2. Claims: 7-19

monohydrates of azithromycin; compositions containing them; and processes for their preparation (other than the process used in subject 1).

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 00/05948

Patent document cited in search report	Publication date	Patent family member(s)		Publication date	
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